

# Effective visible light photodegradation of ortho and para- nitrophenols using BiVO<sub>4</sub>

A. M. Umabala

**Abstract**— T Photocatalytic degradation of ortho- and para nitrophenols was investigated over BiVO<sub>4</sub> under visible light irradiation. BiVO<sub>4</sub> degraded ortho nitrophenol completely in 210 min. Rate of degradation enhanced in presence of H<sub>2</sub>O<sub>2</sub> and complete degradation was achieved in 120min. Unlike ortho nitrophenol, para nitrophenol showed less photolysis and complete degradation of para nitrophenol was achieved for 120min of irradiation in presence of BiVO<sub>4</sub> and H<sub>2</sub>O<sub>2</sub>.

**Index Terms**—A2-nitrophenol, 4-nitrophenol, BiVO<sub>4</sub>, photocatalytic degradation.

## I. INTRODUCTION

Phenols and phenolic compounds are common pollutants of aquatic systems. For instance, nitro phenols are common compounds detected in agricultural waste. Similarly, para nitro phenol used in the synthesis of dyes, pharmaceuticals, pesticides, herbicides and explosives is a common constituent in the effluents from industries involved in manufacturing these chemicals. Besides being carcinogenic and mutagenic, para nitro phenol is toxic even to plants, animals and microorganisms. Though different methods have been employed for the remediation of phenols in terms of solvent extraction, adsorption, membrane separation and chlorination, these methods have some inherent drawbacks as they generate secondary pollution due to phase transfer of pollutants. During the last two decades, attention has been focused on the semiconductor mediated heterogeneous photocatalysis for remediation of nitro phenols using different advanced oxidation processes as discussed in different reviews [1-4].

Giuseppe Marci et al [5] reported degradation of 4-nitro phenol (4-NP) using polycrystalline ZnO/TiO<sub>2</sub> under U.V irradiation for 150min and continuous oxygen bubbling. Di Paola et al [6] studied photocatalytic activity of TiO<sub>2</sub> impregnated with various transition metal ions for 4-NP degradation using 150min of U.V. irradiation. Kashif Naeem and Feng Ouyang [7] reported 35% photocatalytic degradation of 4-NP under U.V irradiation for 120min over TiO<sub>2</sub> dispersed on active carbon, ZSM-5, SiO<sub>2</sub> and rice husk. Lixia Yang et al [8] reported degradation of 4-NP over Cu deposited on TiO<sub>2</sub> heterojunction in 250min under solar light. Hasan Ilyas et al [9] reported 80% degradation of 4-NP over Ag-TiO<sub>2</sub> particles under U.V irradiation for 60min. Wan-Jun Sun et al [10] reported 90% degradation of 4-NP under visible light over Cu porphyrin-TiO<sub>2</sub> +H<sub>2</sub>O<sub>2</sub>. Rahmatollah Rahimi and co workers [11] studied photodegradation of 4-NP over N, S codoped TiO<sub>2</sub> under visible light irradiation for 180min. Hond Ben Ybt Suida and Bassem Jamoussi [12] reported 98%

degradation of 4-NP over TiO<sub>2</sub>-Zinc phthalocyanin under solar light and oxygen bubbling for 60min. Lu Pan et al [13] reported 90% degradation of 4-NP in 180min over CuCr<sub>2</sub>O<sub>4</sub> and H<sub>2</sub>O<sub>2</sub> under visible light irradiation. Safa et al [14] reported 98% degradation of 4-NP in 180min over ZnO-nano flowers under U.V irradiation. Shafiqul Islam et al [15] reported 90% degradation of 4-NP in 120min over TiO<sub>2</sub>+H<sub>2</sub>O<sub>2</sub> under U.V irradiation and observed that addition of Cu<sup>2+</sup> enhances the percent degradation but excess of Cu<sup>2+</sup> decreases the degradation. Sugiyama et al [16] reported that the degradation capability of ZnO particles towards 4-NP under solar radiation was superior to U.V light irradiation. Zhigang Xiong and coworkers [17] reported 90 to 100% degradation of 4-NP in 300min over Au and Pt-TiO<sub>2</sub> composites under visible light. Si Zhan Wu and coworkers [18] reported 90% degradation of 4-NP in 360min over graphitic carbon-nitride (g-C<sub>3</sub>N<sub>4</sub>). Hong Xu Guo et al [19] reported 90% degradation of 4-NP over Zn<sub>3</sub>(VO<sub>4</sub>)<sub>2</sub> under visible light. Radwa Elsalamony and Dalia Abd El- Hafiza [20] reported degradation of 4-NP in 150min over Cu-TiO<sub>2</sub> under U.V irradiation. Nguyen Quang Long et al [21] reported degradation of 4-NP over Fe<sub>3</sub>O<sub>4</sub> -N-doped TiO<sub>2</sub> under visible light with continuous air bubbling. Hyun-Gyu Lee et al [22] reported degradation of 4-NP under visible light using TiO<sub>2</sub>-graphene-palladium nanowires. Khadija Eddanani et al [23] reported degradation of 4-NP over Li<sub>0.5</sub>M<sub>0.5</sub>Ti<sub>2</sub>(PO<sub>4</sub>)<sub>3</sub> (M=Ni, Co and Mn) + H<sub>2</sub>O<sub>2</sub> under visible light. Jing Zhang et al [24] reported 80% degradation of 4-NP in 180min over TiO<sub>2</sub>-Sn-Porphyrin nanoparticles under visible light irradiation. Suranjan Sikdar, and Coworkers [25] studied degradation of 4-NP over M<sub>x</sub>Nb<sub>x</sub>Ti<sub>1-2x</sub>O<sub>2+x/2</sub> M=Cr, Fe; X=0.01-0.2) under U.V irradiation. To our knowledge, so far there are no studies reported on photocatalytic degradation of nitro phenols using BiVO<sub>4</sub> although BiVO<sub>4</sub> is reported to be an excellent visible light responding photocatalyst for the degradation of several dyes [26].

In view of the above, the present work is under taken to investigate the degradation of ortho nitrophenol and para nitrophenols using BiVO<sub>4</sub> as photocatalyst in presence of an external oxidant H<sub>2</sub>O<sub>2</sub> under visible light irradiation without any air/oxygen bubbling.

## II. MATERIALS AND METHODS

### A. Synthesis

Monoclinic BiVO<sub>4</sub> is synthesized using room temperature solid-state metathesis reported elsewhere [27] from this laboratory. A.R.grade BiCl<sub>3</sub> (Loba) was used as precursor along with Na<sub>3</sub>VO<sub>4</sub> (Aldrich). Stoichiometric quantities of reactants in 1:1 molar ratio were weighed and ground thoroughly in an agate mortar for 2hrs in presence of ethanol. The mixture immediately turned to canary yellow in colour. The homogenized mixture was washed with distilled water to

remove NaCl by product and dried at room temperature. The dried sample was used for phase identification and catalytic studies.

### B. Characterizations

Phase identification of the sample was investigated with X-ray diffractometer (PANalytical- X' Pert PRO, Japan) at room temperature, using Nickel filtered Cu-K $\alpha$  radiation ( $\lambda = 1.54059 \text{ \AA}$ ), over a range of  $10^\circ \leq 2\theta \leq 80^\circ$  with a scan rate of  $2^\circ \text{ min}^{-1}$ .

### C. Photocatalytic studies

Photo catalytic activity of BiVO<sub>4</sub> was evaluated in terms of degradation of ortho and para nitro phenols under visible light. 10 mg of the catalyst powder was added into 100ml 2-NP/4-NP aqueous solution (10 mg/L). Before irradiation, the above suspension was magnetically stirred for 30 minutes. The suspension was then exposed to 400 W metal halide lamp; 5ml aliquots were pipetted at periodic time intervals. Progress of decolorization was followed by recording the corresponding absorption spectrum. All the experiments were conducted under ambient conditions. Percent degradation of dye was computed using the relation

$$\% \text{ degradation} = (A_0 - A_t) / A_0 \times 100$$

where  $A_0$  and  $A_t$  are respectively initial absorbance and absorbance at time 't'

## III. RESULTS AND DISCUSSION

BiVO<sub>4</sub> has been reported to exist in three polymorphic modifications namely Tetragonal zircon, Monoclinic scheelite and Tetragonal scheelite. Of these three crystalline modifications, only the monoclinic form of BiVO<sub>4</sub> exhibits visible light induced photocatalytic activity. Fig. 1 depicts X-ray diffraction (XRD) pattern of BiVO<sub>4</sub> sample prepared by solid-state metathesis reaction. All peaks in the XRD pattern could be assigned to monoclinic BiVO<sub>4</sub> of JCPDS File NO 83-1698. Absence of peaks due to any contaminant suggests that the sample obtained is phase pure BiVO<sub>4</sub> of monoclinic structure.

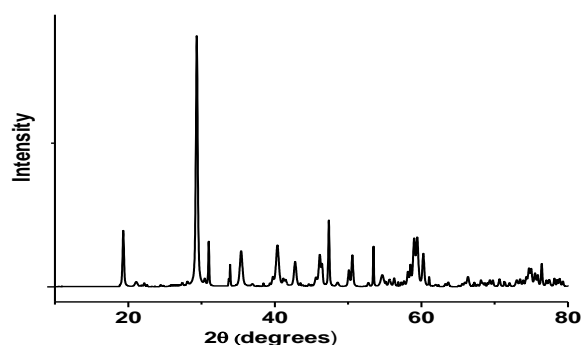


Fig. 1. X-ray diffraction pattern of resultant ground mixture of reactants after washing

In the degradation of dyes with different chromophores, three different approaches were proposed in literature to enhance the photocatalytic quantum efficiency of BiVO<sub>4</sub>. These are (i) synthesis of phase pure monoclinic BiVO<sub>4</sub> crystalline modification (ii) preparing high surface area BiVO<sub>4</sub> through high energy facets and (iii) formation of special architecture

composites such as Bi<sub>2</sub>O<sub>3</sub>-BiVO<sub>4</sub>, Bi<sub>2</sub>S<sub>3</sub>-BiVO<sub>4</sub>, Fe<sub>2</sub>O<sub>3</sub>-BiVO<sub>4</sub> etc. Since synthesis of nano BiVO<sub>4</sub> with high surface area or with special architecture is not cost effective and not viable for large scale industrial applications, addition of external oxidant H<sub>2</sub>O<sub>2</sub> is taken up in this study to enhance the photocatalytic efficiency of m-BiVO<sub>4</sub> because addition of H<sub>2</sub>O<sub>2</sub> has been reported to be beneficial in degradation of several dyes over different photocatalysts [28].

Reports on photocatalytic degradation of ortho nitrophenol (2-NP) are somewhat limited as compared to that of 4-NP. Di Paola et al [29] reported photocatalytic degradation of 2-NP over TiO<sub>2</sub> under U.V irradiation for 240min. Priya and Giridhar [30] reported degradation of 2-NP over TiO<sub>2</sub> under U.V irradiation for 150min. Asha and Sharma [31] reported degradation of 2-NP over Ag-TiO<sub>2</sub> under U.V. irradiation for 360min of irradiation. Jingtao Dai et al [32] reported degradation of 2-NP over TiO<sub>2</sub> nanoparticles synthesized by hydrothermal method using ionic liquids. Aslam et al [33] reported enhanced photocatalytic activity of V<sub>2</sub>O<sub>5</sub>-ZnO composites for the mineralization of 2-NP for 150min of irradiation under sun light. Temporal variation of spectral contours for 2-NP, 2-NP+H<sub>2</sub>O<sub>2</sub>, 2-NP+BiVO<sub>4</sub> and 2-NP+BiVO<sub>4</sub>+H<sub>2</sub>O<sub>2</sub> as a function of irradiation time are shown in Fig 2.

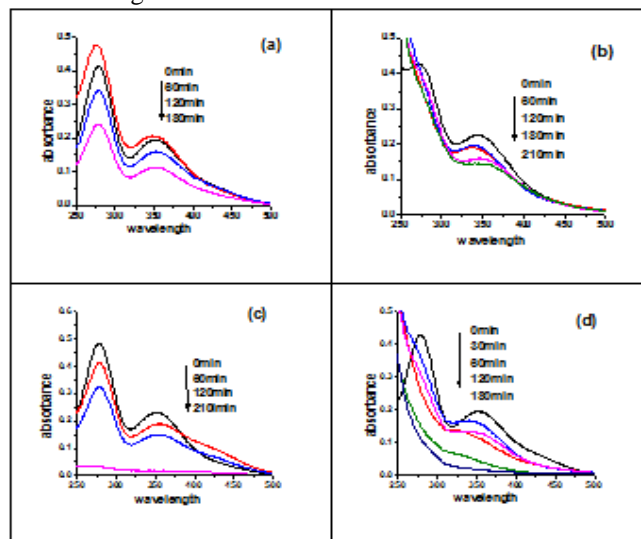


Fig 2. Temporal variations of spectral contours for (a) 2-NP, (b) 2-NP+H<sub>2</sub>O<sub>2</sub>, (c) 2-NP+BiVO<sub>4</sub>, and (d) 2-NP+BiVO<sub>4</sub>+H<sub>2</sub>O<sub>2</sub> as a function of irradiation time.

From the spectra it can be seen that 2-NP exhibits two absorption peaks at  $\lambda = 275$  and  $350 \text{ nm}$  and undergoes slow photodegradation to an extent of  $\sim 42\%$  for irradiation of 180min (Fig 2(a)). In presence of H<sub>2</sub>O<sub>2</sub>, 2-NP shows photodegradation to an extent of  $\sim 36\%$  for irradiation of 210min (Fig 2(b)). However, in presence of BiVO<sub>4</sub> the intensities of both peaks become zero indicating complete degradation (Fig 2(c)) for 210min of irradiation. The rate of photodegradation of 2-NP is found to be expedited in presence of H<sub>2</sub>O<sub>2</sub> since 100% degradation is achieved in 180 min (Fig 2(d)). Fig. 3 shows variation of spectral intensities of 2-NP as a function of irradiation time for 20, 30 and 50mg of catalyst. From the figure it can be seen that 20mg of BiVO<sub>4</sub> is the optimum amount of catalyst required for complete degradation in 120min.

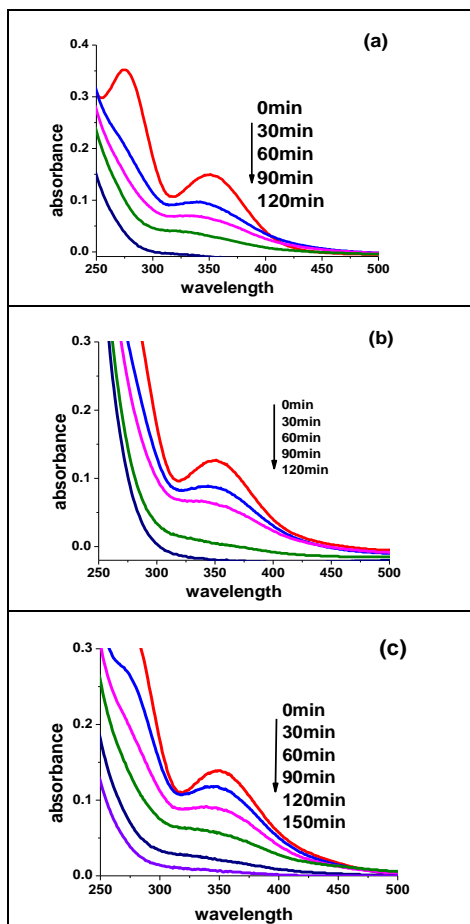


Fig. 3. Temporal variation of spectral changes of 2-NP with (a) 20mg, (b) 30mg and (c) 50mg of photocatalyst in presence of  $H_2O_2$  as a function of irradiation time

The above data clearly indicates that 2-NP can be successfully degraded over  $BiVO_4$  in presence of  $H_2O_2$ . Fig. 4 depicts temporal variation of spectral contours for 4-NP, 4-NP+ $H_2O_2$ , 4-NP+ $BiVO_4$  and 4-NP+ $BiVO_4$ + $H_2O_2$  as a function of irradiation time.

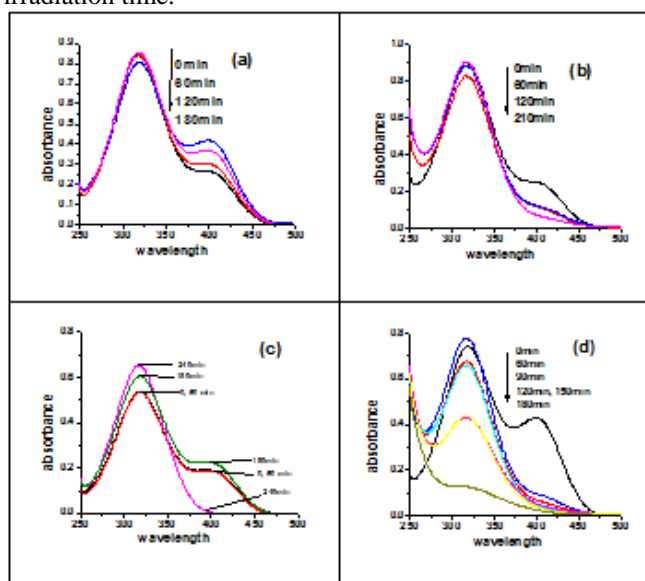


Fig 4. Temporal variation of spectral contours for (a) 4-NP, (b) 4-NP +  $H_2O_2$ , (c) 4-NP +  $BiVO_4$ , and (d) 4-NP +  $BiVO_4$  +  $H_2O_2$  as a function of irradiation time.

The spectra in Fig 4 (a) indicates characteristic absorption for 4-NP at 310 along with a shoulder at 400 nm. From the figure

it can be seen that both 4-NP and 4-NP+ $H_2O_2$  undergo very less photodegradation with progressive irradiation up to 210min (Fig 4 a and b). In contrast to 2-NP, degradation of 4-NP in presence of only  $BiVO_4$  is not observed (Fig.4(c)). However, in presence of  $H_2O_2$  and  $BiVO_4$ , intensities of both peaks decrease to zero indicating near complete degradation of 4-NP for 150min of irradiation (Fig 4(d)). Fig. 5 depicts variation of spectral intensities as a function of irradiation time for 4-NP with 20, 30 and 50mg of photocatalyst keeping the concentrations of 4-NP as well as  $H_2O_2$  unchanged.

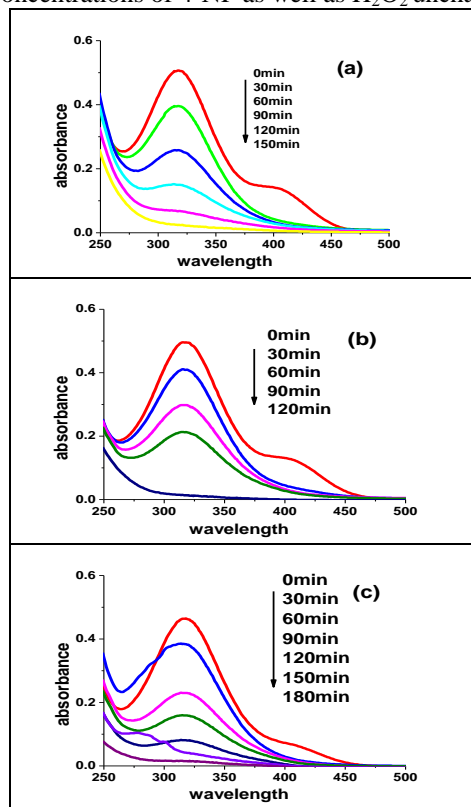
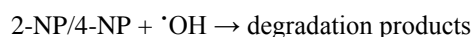
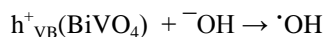
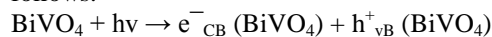


Fig. 5. Temporal variation of spectral changes of 4-NP for (a) 20mg, (b) 30mg and (c) 50mg of photocatalyst in presence of  $H_2O_2$  as a function of irradiation time

From the figure it can be seen that 30mg of  $BiVO_4$  is the optimum amount of catalyst required for irradiation of 120min. Based on the experimental data, the photocatalytic degradation mechanism in presence of  $H_2O_2$  can be given as follows.



The above results show that  $BiVO_4$  can be used as a successful photocatalyst for complete degradation of both ortho and para-nitrophenol in the visible region in presence of  $H_2O_2$  without any  $O_2$  or air bubbling.

## CONCLUSIONS

Complete photocatalytic degradation of ortho and para nitrophenols was successfully achieved by photocatalysis over  $BiVO_4$  in presence of  $H_2O_2$  under visible light irradiation.  $BiVO_4$  effected photocatalytic degradation of ortho nitrophenol even in the absence of  $H_2O_2$  and the

degradation rate increased in presence of H<sub>2</sub>O<sub>2</sub>. BiVO<sub>4</sub> degraded para nitrophenol completely only in presence of H<sub>2</sub>O<sub>2</sub>.

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**A. M. Umabala**, Associate Professor, Dept. of Inorganic & Analytical Chemistry, Andhra University, Visakhapatnam, India-530 003.